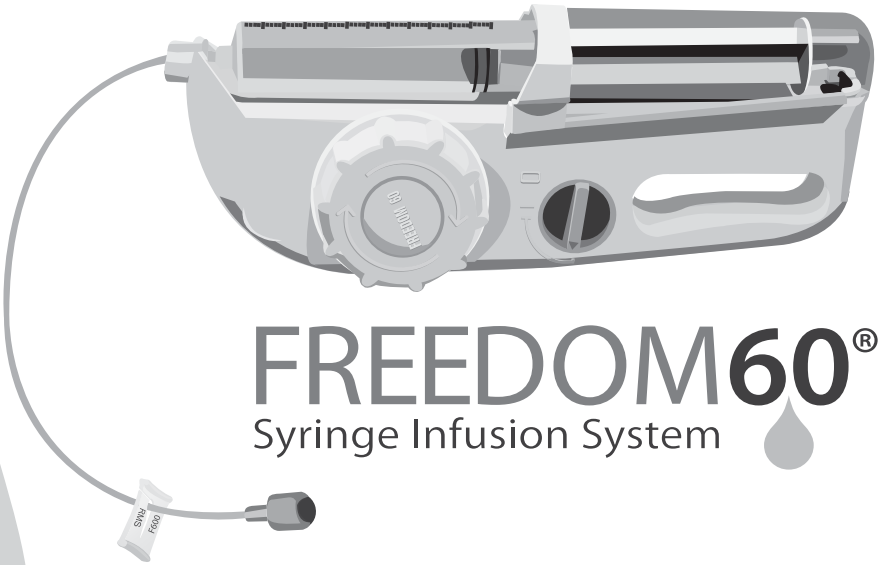

FREEDOM60®

Syringe Infusion System

Instructions For Use

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Introduction:
The FREEDOM60® Syringe Infusion System is portable and easy to use, requiring no batteries or electric. There are only two operating knobs and special precision tubing sets are used to control the flow rate.

The FREEDOM60® operates at a constant safe pressure of 13.5psi. The constant pressure developed in the FREEDOM60® automatically decreases the flow rate if there is an increase in resistance during the delivery. The system will find equilibrium between the increasing resistance and flow rate. It provides constant flow which tends to inhibit clots, and holds full pressure after an infusion is complete to prevent blood or drug backflow. The FREEDOM60® also eliminates concerns of a bolus, overflow, overdose or runaway infusion.

Indications for Use:
The FREEDOM60® Syringe Infusion System is indicated for the infusion of prescribed liquid medicines.

Contraindications:
The FREEDOM60® is not intended for the delivery of blood, critical* or life-sustaining medications, or for the infusion of insulin.

*Critical may be defined as medication requiring greater accuracy of delivery, such as CNS opiate depressants which in certain countries, e.g. UK, may be limited by local regulations. CONSULT LOCAL REGULATORY AUTHORITIES.

Caution:

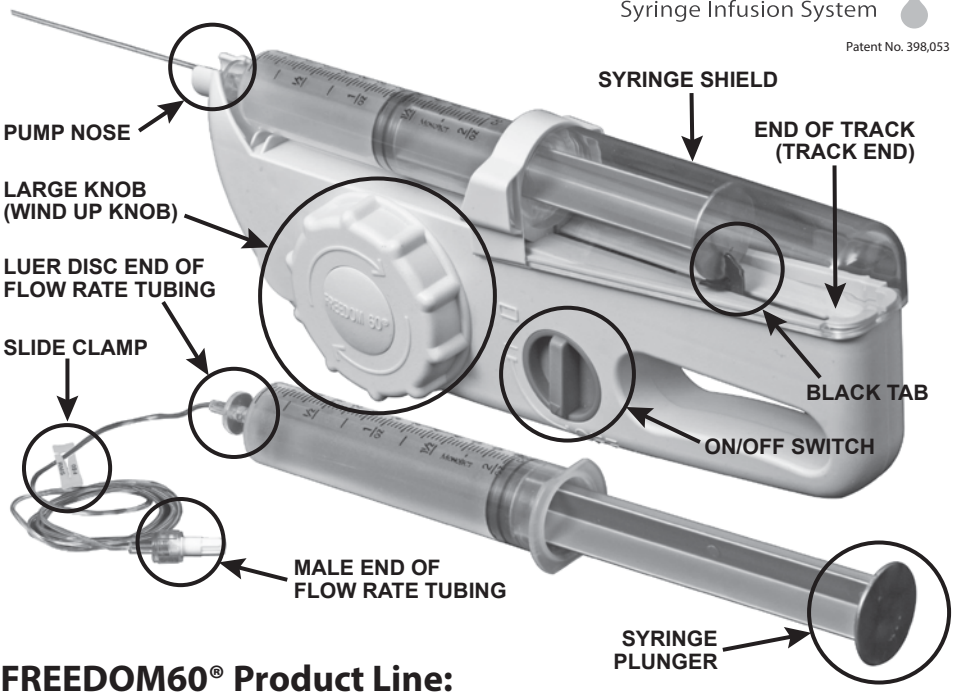
- Use the FREEDOM60® Syringe Infusion System only for the patient for whom the device is prescribed and only for its intended use.
- Do not connect an extension tubing or needle set directly to the syringe. An appropriate flow restrictive tubing with a disc luer is required for use with FREEDOM60®.
- Use only recommended 60ml syringes with the FREEDOM60®. Do not use Terumo® syringes.
- Do not use syringes smaller than 60ml. The use of a smaller syringe may cause high pressures unsafe for the patient.
- Before use, carefully inspect the tubing set package. Do not use the tubing set if the package is opened or damaged.
- Do not re-sterilize tubing sets.
- Overuse of the slide clamp or storing tubing sets with the slide clamp engaged for long periods of time* may damage the tubing and affect the infusion rate.
- The black tab that pushes on the syringe plunger operates under high force. Do not place fingers on the black tab or inside the syringe shield at any time. Do not attempt to interfere with the movement of the black tab at any time.
- Carefully inspect the FREEDOM60® Syringe Infusion Pump before use. Verify its condition and test. If repeated testing indicates the pump is not operating properly or at the appropriate rate of flow, immediately discontinue use.
- Do not attempt to open the pump housing or remove the syringe shield. Do not operate if the syringe shield has been removed.
- Do not attempt to remove the syringe or disconnect the tubing set without first turning the pump OFF and fully winding the large knob clockwise until the black tab has reached the end of its track.
- The FREEDOM60® Syringe Infusion System does not have an alarm, therefore no alarm will sound if an interruption to flow occurs. There is no display of infusion status. The pump is not suitable for use with medication where delay or under-infusion could result in serious injury.
- Discontinue use of a pump that has been damaged, exposed to severe impact, or which has failed to test properly.
- Discontinue use of a pump that has been submerged in fluid. If any fluid is allowed to enter the pump, it should be replaced immediately.
- Do not autoclave the FREEDOM60®. It will melt the ABS plastic and damage the pump.
- Federal law (USA) restricts this device to sale by or on order of a physician.
- Priming and shipping of tubing sets, packaged at temperatures below freezing, is not recommended. This may damage the tubing.

*For example, generally greater than about 2 hours.

FREEDOM60® Pump Diagram:

FREEDOM60®
Syringe Infusion System

Patent No. 398,053



FREEDOM60® Product Line:

Each FREEDOM60® Syringe Infusion System includes a travel pouch and user manual.

Product	Part #
FREEDOM60® Syringe Infusion Pump	F10050
Travel Pouch	F10090

RMS Precision Flow Rate Tubing™ (case of 50)

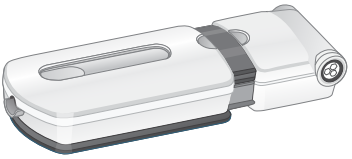
Part #	Flow Rate	Part #	Flow Rate
F0.5	0.5ml/hr[1]	F60	60ml/hr[2]
F1	1ml/hr[1]	F120	120ml/hr[2]
F2	2ml/hr[1]	F180	180ml/hr[3]
F3	3ml/hr[1]	F275	275ml/hr[3]
F3.8	3.8ml/hr[1]	F420	420ml/hr[3]
F5	5ml/hr[1]	F500	500ml/hr[3]
F8	8ml/hr[1]	F600	600ml/hr[3]
F10	10ml/hr[1]	F900	900ml/hr[3]
F15	15ml/hr[1]	F1200	1200ml/hr[3]
F30	30ml/hr[2]	F2400	2400ml/hr[3]
F45	45ml/hr[2]		

precision
FLOW RATE TUBING™

[1] Sterile water for injection (SWFI) direct flow rates.
[2] Adjusted for antibiotic use, for actual flow rates see examples on pages 10-11.
[3] SWFI flow rates will be reduced from stated values. Typically used for viscous drugs.

For Smaller Volume Infusions:

The FREEDOM60® is designed to accommodate 60ml syringes. For smaller volumes, the FreedomEdge™ accommodates both 20ml and 30ml syringes.



FreedomEdge™
Syringe Infusion System

Part #: F10020

Syringes for use with the FREEDOM60®:

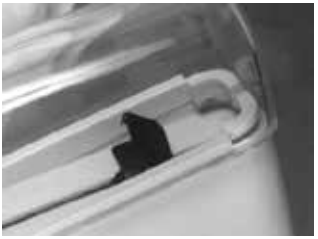
- Becton Dickinson & Co. BD® Luer-Lok® 60ml (US Reference #309653; EU Reference #300865)
- Sherwood Medical Monoject® 60ml (Reference #8881-560125)

Testing the FREEDOM60® Syringe Infusion Pump:

1. Examine the inside of the syringe shield and ensure it is free of debris or contamination.
2. Make sure that the pump's on/off switch is in the OFF position and that the black tab within the syringe shield is at the end of its track. If the black tab is not at the end of its track, fully wind the large knob clockwise.
3. Turn the pump ON (–) and watch that the tab moves smoothly along the full length of its track; listen for the following sounds:
    - a 'click' as the pump is turned ON (–)
    - a 'whirring' sound as the black tab moves forward
    - a 'click' as the tab reaches the end of its travel
4. Test to make sure the syringe tension tab (the one-inch long tab located at the entry of the syringe shield) operates freely by adjusting it up and down with your finger.

Syringe Loading and Removal Instructions:

To load a syringe:



Make sure that the pump's on/off switch is in the OFF position and that the black tab within the syringe shield is at the end of its track. If the black tab is not at the end of its track, wind the large knob clockwise.

With the tubing attached and gradations facing up, load the pre-filled syringe, plunger first, into the syringe shield. Make sure the tubing luer disc is fully seated in the pump's nose.

To remove a syringe:

After the infusion is complete, turn the pump OFF and wind the large knob clockwise until the black tab is at the end of its track. Remove the empty syringe by gently pushing it back to disengage the nose, then lifting it up out of the pump.

Note: You will never need to use force to load or remove a syringe. If you're having trouble loading or removing, stop and make sure the black tab is at the end of its track.

Selected Flow Rates vs Time:

Note: For assistance in determining which flow rate tubing set to use, please contact RMS Medical Products at 800-624-9600.

Flow Rate vs Time Chart

Syringe Volume	1ml/hr	2ml/hr	30ml/hr	45ml/hr	60ml/hr	120ml/hr
5	5 hrs.	2 hrs. 30 min.	10 min.	6 min. 42 sec.	5 min.	2 min. 30 sec.
10	10 hrs.	5 hrs.	20 min.	13 min. 18 sec.	10 min.	5 min.
15	15 hrs.	7 hrs. 30 min.	30 min.	20 min.	15 min.	7 min. 30 sec.
20	20 hrs.	10 hrs.	40 min.	26 min. 42 sec.	20 min.	10 min.
25	25 hrs.	12 hrs. 30 min.	50 min.	33 min. 18 sec.	25 min.	12 min. 30 sec.
30	30 hrs.	15 hrs.	60 min.	40 min.	30 min.	15 min.
35	35 hrs.	17 hrs. 30 min.	70 min.	46 min. 42 sec.	35 min.	17 min. 30 sec.
40	40 hrs.	20 hrs.	80 min.	53 min. 18 sec.	40 min.	20 min.
45	45 hrs.	22 hrs. 30 min.	90 min.	60 min.	45 min.	22 min. 30 sec.
50	50 hrs.	25 hrs.	100 min.	66 min. 42 sec.	50 min.	25 min.
55	55 hrs.	27 hrs. 30 min.	110 min.	73 min. 18 sec.	55 min.	27 min. 30 sec.
60	60 hrs.	30 hrs.	120 min.	80 min.	60 min.	30 min.

Starting the Infusion:

Before loading the syringe verify the following:

- Make sure that the pump's on/off switch is in the OFF position.
- Make sure that the black tab inside the syringe shield is at the end of its track. If the black tab is not at the end of its track, wind the large knob clockwise.
- Verify that you have the correct rate controlled tubing set.

1. Using sterile technique, remove the sterile cap from the luer disc end of the rate controlled tubing set and connect the tubing to the pre-filled syringe.
2. With syringe gradations facing up, insert the syringe, plunger first into the syringe shield. Make sure the luer disc is fully seated in the pump's nose.

Note: You will never need to use force to load or remove a syringe. If you are having trouble loading/removing, make sure the pump is in the OFF position and that the black tab is at the end of its track. If the black tab is not at the end of its track, fully wind the large knob clockwise.

3. Using sterile technique, prime the tubing with the pump by loosening the cap on the male end and turning the pump ON (–). You will immediately hear a "whirring" sound as the black tab searches for the syringe plunger. When the tab finds the plunger, it will "click" indicating that that the tab is now applying pressure onto the plunger. Medication is now being pumped through the tubing set. When all air and air bubbles are gone and medication starts to drip, tighten the cap.

NOTE: *Tubing that is pre-primed for antibiotic use and shipped at temperatures below freezing could be damaged.*

- Test the tubing: While the pump is in the ON (–) position, use sterile procedures to disconnect the rate controlled tubing set from the indwelling needle or catheter; check for medication drip. If medication does not drip, replace the tubing as it may be blocked or damaged. If medication does drip from the rate controlled tubing, then it's most likely a problem with the indwelling needle, catheter, or any fluid path accessories such as a clave connector or needle-free adapter.

Slow flow

- If a slide clamp is used for an extensive period of time, it can damage the tubing and affect the flow rate. Try using another tubing set and measure the flow. For 60ml/hr tubing, the syringe plunger should move 10ml in 10 minutes (1ml/min). For 120ml/hr tubing, the plunger should move 10ml in 5 minutes (2ml/min).

Continuous flow even when the pump is turned OFF

- This is a normal function of the pump. The pump is designed to maintain pressure during and after the infusion to prevent blood/drug backflow.
- To stop the flow turn the pump OFF and wind the large knob clockwise so the black tab moves away from the syringe. This will release pressure on the plunger.
- You can also use the slide clamp. We recommend using the slide clamp to immediately stop the flow only in the case of an emergency. Overuse can damage the tubing.

Medication (5ml or less) left in the syringe

- Verify that you are using the proper BD® or Monoject® brand 60ml syringe. Do not use Terumo syringes.

Care and Maintenance:

The FREEDOM60® does not require any preventative maintenance. The FREEDOM60® works as a system, which means the tubing determines the flow rate, not the pump; therefore our pump needs no calibration. If you choose the correct tubing set the proper flow rate will be achieved. Our tubing sets are statistically verified during manufacture and will deliver the appropriate flow rate under controlled conditions.

Cleaning

Clean only those areas that are exposed and external. No attempt should be made to clean any part of the pump that is not easily accessible. Discontinue use of a pump that has been internally exposed to or immersed in fluid.

Wipe the surface with warm water and detergent or use any surface disinfectant compatible with acetyl-butyl-styrene (ABS) and polycarbonate (PC) plastics, such as household bleach or hydrogen peroxide. Avoid the use of alcohol or alcohol containing compounds, as these tend to make ABS plastic brittle. Wipe again with clean water to rinse.

- Test the tubing: While the pump is in the ON (–) position, use sterile procedures to disconnect the rate controlled tubing set from the indwelling needle or catheter; check for medication drip. If medication does not drip, replace the tubing as it may be blocked or damaged. If medication does drip from the rate controlled tubing, then it's most likely a problem with the indwelling needle, catheter, or any fluid path accessories such as a clave connector or needle-free adapter.

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Wipe the surface with warm water and detergent or use any surface disinfectant compatible with acetyl-butyl-styrene (ABS) and polycarbonate (PC) plastics, such as household bleach or hydrogen peroxide. Avoid the use of alcohol or alcohol containing compounds, as these tend to make ABS plastic brittle. Wipe again with clean water to rinse.

If absolutely necessary, you may clean the inside of the syringe shield by using a long swab or bottlebrush and any cleaning agent compatible with ABS and PC plastics. Alcohol is not recommended.

Storage:
The FREEDOM60® pump is recommended to be stored in a cool, dry place and packaged tubing sets at room temperature (approximately 16-30°C or 61-86°F).

Testing Flow Accuracy (if required by your local protocol):

1. Fill a new syringe with 60ml of sterile water
2. Remove all air from the syringe
3. Attach a sterile RMS Precision Flow Rate Tubing™ set to the syringe
4. Remove all air from the tubing set
5. Load the syringe into the pump and turn the pump ON (–)
6. Monitor the syringe readings and elapsed time to derive an approximate flow rate
7. Compare your test results to the range of test rates listed in the table below:

Labeled Flow Rate	Bench-Rated Flow Rate	Test Range
60ml/hr	72ml/hr	60-84ml/hr
120ml/hr	134 ml/hr	115-153ml/hr

7a. **Bench-Rated Flow Rate:** The FREEDOM60® infusion pump design accounts for the effects of standard clinical conditions on flow rate performance. Under bench test conditions, a 60ml/hr labeled tubing set is designed to generate a nominal infusion rate of 72ml/hr. A 120ml/hr labeled tubing set will generate a nominal bench test rate of 134ml/hr. The FREEDOM60® generates nominal bench test rates higher than the labeled rate accounting for the following standardized application criteria that affects actual delivery rates under normal clinical circumstances.

	60 ml/hr Tubing	120 ml/hr Tubing
Bench Test Rate	72ml/hr	134ml/hr
Less Clinical Effects		
Catheter Gauge (20G PICC)	-7ml/hr	-8ml/hr
Fluid Viscosity	-2ml/hr	-3ml/hr
Venous Pressure	-3ml/hr	-3ml/hr
Label Flow Rate	60ml/hr	120ml/hr

7b. **Test Range:** To assure consistent test results, keep pump and tubing at the same approximate horizontal plane and monitor flow for a minimum of 20 minutes. The FREEDOM60® system is factory rated to deliver infusions under strict test conditions over a large number of pumps tested within 7% of nominal with a 95% statistical confidence interval. Under varying bench test and fluid conditions this range can be expected to vary approximately 15% of nominal. For more accurate monitoring, use a stopwatch and finely graduated burette. FREEDOM60® pump testing is based on ANSI/AAMI National Standard, ID 26-1992, Infusion Devices, August 24, 1992.

If test results in the range indicated cannot be approximated under bench testing conditions, factory refurbishment and full certification testing is available from the manufacturer.

References:

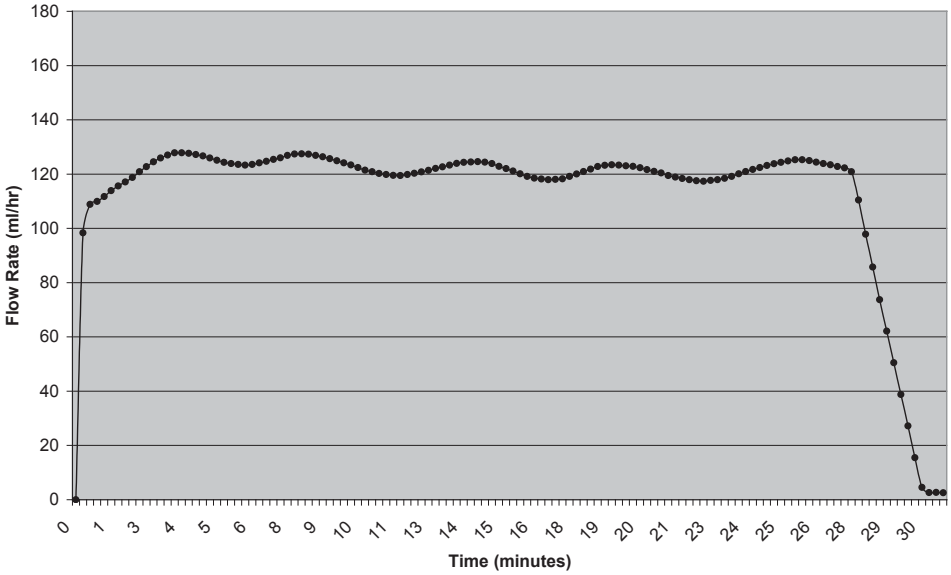
1. Stuhmeier, Mainzer B. MD; Aspects of pressure build-up in the use of electronic infusion devices. II. Need for a pressure limit. Anasth Intensivther Notfallmed (1987 Aug.) 22(4): 185-190.
2. Anasth Intensivther Notfallmed (1987 Aug.) 22(4): 181-184. ANSI/AAMI National Standard, ID 26-1992, Infusion Devices, August 24, 1992.

FREEDOM60® Flow Profile:

The FREEDOM60® flow profile shows that the flow rate is consistent throughout the delivery of medication.

FREEDOM60® Flow Rate vs. Time

Fluid: H_2O • Fluid Volume: 60ml • Tubing Measured: F120 – 120ml/hr



Technical Specifications:

System

Reservoir volume:	60ml (maximum)
Residual volume:	<0.4ml
Flow rate accuracy:	± 7%*
Operating pressure:	15psi (peak) 13.5psi (nominal)
Height sensitivity:	± 3% per 12" (30cm)
Constant force mechanism:	± 3%

*Flow rate data recorded at 22°C using 60ml of 0.9% NS. An overall accuracy of ±7% is expected at these values. At higher temperatures, which results in a decrease in viscosity, a higher flow rate is anticipated to occur. The flow rate variation due to changes in temperature is approximately linear and would vary from -20% at 14°C to +20% at 30°C. Fluids considerably more viscous than 0.9% NS are expected to slow the flow rate and consequently may result in longer infusion times. More viscous fluids may be tested before patient use by following the bench test procedures described in this manual.

Pump

Weight:	14oz (0.4kg)
Length:	12" (304mm)
Width:	4.5" (114mm)
Height:	1.6" (41mm)

RMS Precision Flow Rate Tubing™

Length: 6" to 72" (152mm-1829mm)

RMS Precision Flow Rate Tubing™ / Residual Volume (ml)

F0.5	0.09	F60	0.14
F1	0.08	F120	0.16
F2	0.10	F180	0.13
F3	0.09	F275	0.11
F3.8	0.09	F420	0.10
F5	0.08	F500	0.09
F8	0.08	F600	0.09
F10	0.14	F900	0.08
F15	0.11	F1200	0.13
F30	0.13	F2400	0.15
F45	0.11		

precision
FLOW RATE TUBING™

Warranty Information:

Limited Warranty: RMS Medical Products/Repro Med Systems, Inc. ("Manufacturer") warrants the infusion pump to be free from defects in materials and workmanship under normal use. Warranty is limited to Original Purchaser and covers the FREEDOM60® for a period of two years from the purchase date. This warranty is not valid for any damage caused by the use of non-RMS products. The "Original Purchaser" is the person purchasing the infusion pump from the Manufacturer or Manufacturer's Representative. Warranty does not extend to subsequent purchasers. Subject to the conditions of and upon compliance with the procedures set forth in this limited warranty, the Manufacturer will repair or replace, at its option, any infusion pump, or part thereof, which has been actually received by the Manufacturer or Manufacturer's Representative within the two-year warranty period, and which examination discloses, to the Manufacturer's satisfaction, that the product is defective. Replacement product and parts are warranted only for the remaining portion of the original two-year warranty period.

RMS rigorously tests the FREEDOM60® using RMS accessories to ensure that the FREEDOM60® operates in accordance with published specification standards. If non-RMS accessories are used in conjunction with the FREEDOM60®, RMS does not represent that the FREEDOM60® will operate in accordance with published specification standards. The FREEDOM60® warranty does not cover third-party products or accessories.

The following conditions, procedures, and limitations apply to the Manufacturer's obligations under this warranty:

- **Parties Covered by this Warranty:** This warranty extends only to the Original Purchaser of the infusion pump. This warranty does not extend to subsequent purchasers.
- **Warranty Performance Procedure:** Notice of the defect must be made in writing to Customer Support Department, RMS Medical Products/Repro Med Systems, Inc., 24 Carpenter Road, Chester, NY 10918, USA. Notice to RMS Medical Products/Repro Med Systems, Inc. must include the model and serial number, date of purchase, and description of the defect in sufficient detail to facilitate repairs. Authorization must be obtained by the Original Purchaser from the Manufacturer or Manufacturer's Representative prior to returning the product to the Manufacturer. The defective pump must be properly packaged and returned to the Manufacturer, postage-prepaid. Any loss or damage during shipment is at the risk of the Original Purchaser.
- **Conditions of Warranty:** This warranty does not apply to any product, or part thereof, which has been repaired or altered outside of the Manufacturer's facility in a way so as, in Manufacturer's judgment, to affect its stability or reliability, or which has been subjected to misuse, negligence or accident.
- **Limitations and Exclusions:** Repair or replacement of an infusion pump or component part is the EXCLUSIVE remedy offered by the Manufacturer. The following exclusions and limitations shall apply:
    - o No agent, representative, or employee of the Manufacturer has authority to bind the Manufacturer to any representation or warranty, expressed or implied, or to change this limited warranty in any way.
    - o THIS LIMITED WARRANTY IS IN LIEU OF ALL OTHER WARRANTIES, EXPRESS OR IMPLIED, INCLUDING, BUT NOT LIMITED TO, WARRANTIES OF MERCHANTABILITY AND FITNESS FOR A PARTICULAR PURPOSE. THERE ARE NO WARRANTIES THAT EXTEND BEYOND THE DESCRIPTION ON THE FACE HEREOF.
    - o Manufacturer's liability under this Limited Warranty Agreement shall not extend to special, indirect, or consequential damages.
    - o The infusion pump can only be used under the supervision of medical personnel whose skill and judgment determine the suitability of the infusion pump for a particular medical treatment.
    - o All recommendations, information, and descriptive literature supplied by the Manufacturer or its agents are believed to be accurate and reliable, but do not constitute warranties.

This warranty and the rights and obligations hereunder, shall be construed under and governed by the laws of the State of New York, USA.

Definition of symbols:

_	On		Manufacturer
OFF	Off	QTY	Quantity
	Caution	REF	Catalog number
	Consult instructions for use		Do not reuse
EC REP	Authorized representative in the European Community	SN	Serial number
	Use by YYYY-MM-DD or YYYY-MM	STERILE R	Sterilized using irradiation
LOT	Batch code		

RMS™
MEDICAL PRODUCTS

EC REP

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www.rmsmedicalproducts.com

CE 0297

The FREEDOM60® Syringe Infusion System and RMS Precision Flow Rate Tubing™ are compliant with Medical Device Directive 93/42/EEC. RMS Medical Products is ISO 13485 certified.